PATENT Attorney Docket No.: JHU1520-2

In re Application of
Worley and Brakeman
U.S. Serial No.: 09/910,706

Filed: July 20, 2001

Page 3

B. In the Claims

Please cancel claims 1 to 9, 11 to 13, and 24 to 43 without prejudice; amend claims 10, 14, and 17 to 23 as indicated; and add new claims 44 and 45. Upon entry of the present amendment, the status of the claims will be as follows:

- 1-9. (canceled)
- 10. (currently amended) A method of selecting screening assay for identifying a compound that interferes with binding of a synaptic activation protein to a cellular binding protein metabotropic glutamate receptor (mGluR), comprising:
 - a) adding a test compound to a reaction mixture containing comprising
 - (i) an isolated synaptic activation protein having at least 70% sequence identity to a polypeptide having the sequence comprising SEQ ID NO: 2, and
 - (ii) an isolated binding protein to which mGluR comprising SSSL (SEQ ID NO: 10) or SSTL (SEQ ID NO: 11), wherein the synaptic activation protein binds to the mGluR in the absence of the test compound, and
 - b) (iii) means for detecting a change in binding between the synaptic activation protein and the binding protein; measuring binding between the synaptic activation protein and the binding protein; and selecting the compound if the measured binding is greater than or less than binding measured in the absence of the test compound mGluR in the presence of the test compound as compared to the absence of the test compound, thereby identifying a compound that interferes with binding the synaptic activation protein and the mGluR.

11 to 13. (canceled)

PATENT

Attorney Docket No.: JHU1520-2

 In re Application of Worley and Brakeman
 U.S. Serial No.: 09/910,706

Filed: July 20, 2001

Page 4

14. (currently amended) The method of claim [[13]] 10, wherein the mGluR is selected from mGluR5 and or mGluR1α.

- 15. (currently amended) The method of claim 10, wherein the synaptic activation protein is a Homer protein having the sequence comprising SEQ ID NO: 2.
- 16. (previously presented) The method of claim 10, wherein the synaptic activation protein is coated onto a solid phase.
- 17. (previously presented) The method of claim 16, wherein the solid phase is a microtiter plate.
- 18. (currently amended) The method of claim 10, wherein the means for detecting binding is synaptic activation protein comprises a fusion protein comprising glutathione-S-transferase (GST) pulldown.
- 19. (currently amended) The method of claim 10, wherein the means for detecting a change in binding is performed using a co-immunoprecipitation assay.
- 20. (currently amended) The method of claim 10, wherein the mGluR measuring binding comprises a detectable label labeling the binding protein, wherein the labeling is direct labeling or is subsequent addition of a labeled, binding protein specific reagent.
- 21. (currently amended) The method of claim 10 [[20]], wherein the binding protein-specific reagent is detecting a change in binding comprises contacting the reaction mixture with an antibody specific for the mGluR, wherein the antibody is detectably labeled, and measuring

Attorney Docket No.: JHU1520-2

In re Application of Worley and Brakeman

U.S. Serial No.: 09/910,706

Filed: July 20, 2001

Page 5

antibody bound to mGluR in the presence of the test compound as compared to the absence of the test compound.

- 22. (currently amended) The method of claim 20, wherein the labeling detectable label comprises use of an enzyme capable of generating a signal, use of a radiolabeled reagent, use of a fluorescent dye, or use of gold, or biotin.
- 23. (previously presented) The method of claim 22, wherein the radiolabeled reagent is labeled with ¹²⁵I.

24 to 43. (canceled)

- 44. (new) The method of claim 10, wherein detecting a change in binding is performed using a two hybrid assay.
- 45. (new) The method of claim 18, wherein the fusion protein, which comprises the synaptic activation protein and GST, is linked to a solid support.